(b) the composition further comprises a second tumor associated polypeptide or an immunogenic fragment thereof which is a non-RUR-1 antisense cDNA-encoded tumor associated polypeptide or an immunogenic fragment thereof; or

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(c) the cell further comprises a second tumor associated nucleic acid or polypeptide, which is a non-RUR-1 antisense cDNA nucleic acid or tumor associated polypeptide, or an immunogenic fragment thereof.

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 50.(amended) The vaccine composition of claim <u>47</u> [49], wherein the RUR-1 antisense cDNA-encoded tumor associated polypeptide or an immunogenic fragment thereof comprises the amino acid sequence of SEQ ID NO:3.

\$4.(amended) The vaccine composition of [any of] claim[s] 47[-53], further comprising an adjuvant or a pharmaceutically acceptable carrier.

61.(amended) The method of claim 60 wherein the agent is selected from the group consisting of pa nucleic acid molecule comprising SEQ ID NO:1 or a unique fragment thereof; a cytolytic T symphocyte; and an antibody or antibody fragment.

## Remarks

Applicants have amended certain claims to eliminate multiple dependencies and/or clarify claim language. Subject matter added to the claims is supported in the claims as filed and in the specification. No new matter has been added.

Respectfully submitted,

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